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Azithromycin-liposomes as a novel approach for localized therapy of cervicovaginal bacterial infections

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Background: Efficient localized cervicovaginal antibacterial therapy, enabling the delivery of antibiotic to the site of action at lower doses while escaping systemic drug effects and reducing the risk of developing microbial resistance, is attracting considerable attention. Liposomes have been shown to allow sustained drug release into vaginal mucosa and improve delivery of antibiotics to bacterial cells and biofilms. Azithromycin (AZI), a potent broad-spectrum macrolide antibiotic, has not yet been investigated for localized therapy of cervicovaginal infections, although it is administered orally for the treatment of sexually transmitted diseases. Encapsulation of AZI in liposomes could improve its solubility, antibacterial activity, and allow the prolonged drug release in the cervicovaginal tissue, while avoiding systemic side effects.

Purpose: The objective of this study was to develop AZI-liposomes and explore their potentials for treating cervicovaginal infections.

Methods: AZI-liposomes that differed in bilayer elasticity/rigidity and surface charge were prepared and evaluated under simulated cervicovaginal conditions to yield optimized liposomes, which were assessed for antibacterial activity against several planktonic and biofilm-forming *Escherichia coli* strains and intracellular *Chlamydia trachomatis*, *ex vivo* AZI vaginal deposition/penetration, and *in vitro* cytotoxicity toward cervical cells.

Results: Negatively charged liposomes with rigid bilayers (CL-3), propylene glycol liposomes (PGL-2) and deformable propylene glycol liposomes (DPGL-2) were efficient against planktonic *E. coli* ATCC 700928 and K-12. CL-3 was superior for preventing the formation of *E. coli* ATCC 700928 and K-12 biofilms, with IC₅₀ values (concentrations that inhibit biofilm viability by 50%) up to 8-fold lower than those of the control (free AZI). DPGL-2 was the most promising for eradication of already formed *E. coli* biofilms and for treating *C. trachomatis* infections. All AZI-liposomes were biocompatible with cervical cells and improved localization of the drug inside vaginal tissue compared with the control.

Conclusion: The performed studies confirm the potentials of AZI-liposomes for localized cervicovaginal therapy.

Keywords: vaginal drug delivery, biofilm, *Escherichia coli*, *Chlamydia trachomatis*, cervical cells, biocompatibility

Introduction

Bacterial cervicovaginal diseases, such as sexually transmitted *Chlamydia trachomatis* infections, recurrent bacterial vaginosis, and urogenital infections caused by *Escherichia coli*, represent a major public health problem with a substantial and expanding economic burden on the healthcare system. Vaginal *E. coli* strains are

considered to be a reservoir for vaginal and/or endocervical colonization in pregnant women, which may result in progression of urinary tract, intra-amniotic and postpartum infections. Consequently, maternal or fetal complications can occur, including postpartum endometritis, bacteremia or sepsis.¹ Untreated *C. trachomatis* infections may lead to infertility, adverse pregnancy outcomes, fetal loss and an increased risk of HIV transmission.²

The recommended treatment regimens for genital *E. coli* and *C. trachomatis* infections include oral administration of antibiotics.^{3,4} However, the incidence of antibiotic-related side effects and concerns about bacterial resistance make it necessary to consider other modes of tackling cervicovaginal infections, such as those based on localized cervicovaginal therapy.

Application of liposomally encapsulated antibiotics presents a promising strategy for enhanced drug delivery to bacterial cells and biofilms.⁵⁻⁷ Due to their phospholipid composition, liposomes are of particular interest regarding the safety issue and targeting ability.⁸ Tuning the physicochemical properties of liposomes, such as bilayer composition and fluidity, size, surface charge and coating, allows the development of drug-containing liposomes for localized therapy.^{9,10}

Azithromycin (AZI) is a potent, broad-spectrum macrolide antibiotic that is available in dosage forms for oral, ophthalmic and parenteral administration. It is recommended for the treatment of respiratory, skin and soft tissue infections, including sexually transmitted bacterial diseases caused by *C. trachomatis* and *Neisseria gonorrhoeae*.^{11,12} AZI is usually available as a dihydrate (Mw 785) and is characterized by its limited solubility in water (log *P*=3.98).¹³ Incorporation of AZI into liposomes could increase its solubility, favor its cervicovaginal biocompatibility, and enable prolonged AZI release, permitting higher local drug concentrations.

Although AZI-liposomes have been tested against several pathogens,¹⁴⁻¹⁸ to the best of our knowledge, they have not been investigated for localized treatment of vaginal infections, and conventional topical formulations of AZI are not available for vaginal administration.

Therefore, for the first time, we investigated the potential of several types of AZI-liposomes, ie, conventional (CLs), propylene glycol (PGLs) and deformable propylene glycol liposomes (DPGLs), for the treatment of cervicovaginal infections. The different AZI-liposomes were characterized for the physical properties, in vitro release under simulated cervicovaginal conditions, and storage stability. The optimized liposomes were examined for *ex vivo* permeability

on porcine vaginal mucosa, and their in vitro antibacterial activity was tested against several strains of biofilm-forming *E. coli* and intracellular *C. trachomatis*. Finally, the biocompatibility of the different AZI-liposomes was estimated in an in vitro cervical cell model.

To the best of our knowledge, AZI-liposomes have not yet been assessed against *C. trachomatis* and biofilm-forming *E. coli*, and their in vitro compatibility with cervical cells has not yet been examined. Moreover, this study is the first to employ direct quantitative polymerase chain reaction (qPCR) measurements to study the antichlamydial effect of liposomally encapsulated antibiotics. Finally, the use of monoacyl phosphatidylcholine as an edge activator for the preparation of DPGLs represents a novel approach for the development of liposomes with elastic bilayers.

Materials and methods

Materials

Egg phosphatidylcholine (EPC), egg phosphatidylglycerol sodium salt (EPG), hydrogenated soybean phosphatidylcholine (SPC-3) and monoacyl phosphatidylcholine from soybean (SLPC-80) were kindly donated by Lipoid GmbH (Ludwigshafen, Germany). AZI in the form of a dihydrate was a gift from PLIVA Croatia Ltd. (Zagreb, Croatia). Sephadex G-50, urea, glucose, lactic acid, Dulbecco's modified Eagle's medium, Minimum Essential Medium with Earle's salts, L-glutamine, fetal bovine serum and bovine serum albumin were obtained by Sigma-Aldrich (St. Louis, USA). Acetonitrile, ethanol and methanol were of HPLC grade and purchased from BDH Prolabo (Lutterworth, UK). Luria-Bertani (LB) agar and LB broth were obtained from Fisher Scientific (Leicestershire, UK) and Serva Electrophoresis GmbH (Heidelberg, Germany), respectively. Heat-inactivated fetal bovine serum was purchased from Euroclone (Pero, Italy). All other solvents or chemicals used in this research were of analytical grade and obtained from Kemika (Zagreb, Croatia) or Sigma-Aldrich.

A solution of 0.01 M phosphate-buffered saline (PBS, pH 7.5) was prepared and filtered according to Rukavina et al¹⁷. Vaginal fluid simulant (VFS) was made by dissolving 0.222 g of Ca(OH)₂, 1.40 g of KOH, 3.51 g of NaCl, 2 g of lactic acid, 1 g of acetic acid, 0.16 g of glycerol, 0.4 g of urea, 5 g of glucose and 0.018 g of bovine serum albumin in demineralized water (up to 1000 ml) and adjusting the pH to 4.5 by the addition of 0.1 M HCl.¹⁹ Commercially available PBS,

pH 7.3–7.5 (Lonza, Verviers, Belgium), was used in antibacterial assays against *E. coli*.

Preparation of AZI-liposomes

CLs were prepared by a modified proliposome method.²⁰ In brief, phospholipids and the drug (Table 1) were dissolved in warm ethanol (300 mg), followed by the addition of 0.5 ml of PBS, pH 7.5, with continuous magnetic stirring (600 rpm, 60°C) to yield an initial proliposome mixture. This mixture was cooled to room temperature and then transformed into a liposomal dispersion by slow addition of PBS, pH 7.5, to a final volume of 10 ml of liposome dispersion. The dispersion was stirred (600 rpm) for approximately 45 min at room temperature, while in the case of liposomes embedding SPC-3 (denoted as CL-2 and CL-3), the whole procedure was performed at approximately 50°C.

For the preparation of elastic liposomes (PGLs and DPGLs), the selected phospholipids and AZI (Table 1) were dissolved in propylene glycol using a magnetic stirrer (600 rpm, 50°C). Then, 1 ml of PBS, pH 7.5 (heated to the same temperature) was added. The mixture was stirred for approximately 2 min (50°C) and diluted to a liposomal dispersion by dropwise addition of 8 ml of PBS, pH 7.5, with continuous magnetic stirring (45 min).

The corresponding empty liposomes were prepared under the same conditions, omitting AZI, and were used as controls in in vitro antibacterial and biocompatibility studies.

All the liposomal dispersions were hand-extruded (LiposoFast Basic, Avestin, Canada) through 400 nm polycarbonate membranes (3 cycles) at room temperature and stored at 4°C. The extrusion of CL-2 and CL-3 was performed at 50°C.

Size and zeta potential measurements

The mean diameters, polydispersity indexes (PDIs) and zeta potentials of all the prepared liposomes were determined on a Zetasizer 3000HS (Malvern Instruments, Malvern, UK) at 25°C using a scattering angle of 90° (size measurements) and a capillary cell with an optical modulator at 1,000 Hz (zeta potential measurements). Prior to the measurements, the instrument was calibrated with Malvern's size and zeta potential standards. The liposomal samples were prepared by diluting the liposomal dispersions with PBS, pH 7.5, to achieve appropriate count rates.²¹ The measurements were also performed in VFS, pH 4.5, at 37°C and 25°C.

Determination of liposome bilayer elasticity/rigidity

The elasticities/rigidities of the different AZI-liposome bilayers were determined using a previously reported method.²² In brief, liposomes were extruded through a 100 nm pore-sized membrane (r_p) for 5 min at an external pressure of 2.5 bar. The mass of the extruded liposomes (J) and the mean diameter of the liposomes after extrusion (r_v) were determined. The degree of bilayer elasticity (E) was calculated by applying the following equation:²¹

$$E = J \cdot (r_v/r_p)^2$$

Encapsulation efficiency determination and quantification of AZI

The encapsulation of AZI in the different liposomes was determined after separation of unencapsulated AZI by a minicoloumn centrifugation method.²³ In brief, minicoloumns filled with Sephadex G-50 gel were dried by

Table 1 Composition of liposomes

Liposomes type	Code	EPC (mg)	EPG (mg)	SPC-3 (mg)	SLPC-80 (mg)	AZI (mg)	PG (g)	PBS, pH 7.5 (ml)
CLs	CL-1	190	10	-	-	30	-	10
	CL-2	150	-	50	-	30	-	10
	CL-3	140	10	50	-	30	-	10
PGLs*	PGL-1	200	-	-	-	30	1	9
	PGL-2	190	10	-	-	30	1	9
DPGLs*	DPGL-1	170	-	-	30	30	1	9
	DPGL-2	160	10	-	30	30	1	9

Notes: The final volume of liposome dispersion was 10 ml. *Elastic liposomes.

Abbreviations: AZI, azithromycin; CLs, conventional liposomes; DPGLs, deformable propylene glycol liposomes; EPC, egg phosphatidylcholine; EPG, egg phosphatidylglycerol sodium salt; PBS, phosphate-buffered saline; PG, propylene glycol; PGLs, propylene glycol liposomes; SLPC-80, soybean monoacyl phosphatidylcholine; SPC-3, hydrogenated soybean phosphatidylcholine.

centrifugation (2,000 rpm, 3 min). Then, 0.4 ml of liposome dispersions were cautiously applied to the top of the gels, followed by centrifugation under the same conditions to obtain AZI-liposomes in the centrifuge tubes, which were saved for assessment. Subsequently, 0.5 ml of PBS, pH 7.5, was added to each minicolumn, which was again centrifuged, and the eluates were collected. The last procedure, including rinsing the minicolumns with PBS, was repeated until all unencapsulated AZI was removed.

Quantification of AZI, both unencapsulated and liposomally encapsulated, was performed by HPLC under the previously described conditions.¹⁷

The encapsulation of AZI was expressed as the encapsulation efficiency (%), ie, the percentage of encapsulated AZI compared to the original drug concentration and the amount of encapsulated drug per mass of phospholipids used.

In vitro release studies

The release of AZI from the different liposomes, which were previously separated from the unencapsulated drug, was assessed under conditions simulating vaginal environment using the so-called bag dialysis method. AZI-liposomes (corresponding to 1 mg of AZI) were placed in a dialysis bag (Mw cut off 12–14,000 Da, Medicell Membranes Ltd., London, UK) and dialyzed against 25 ml of VFS (pH 4.5) with constant stirring (50 rpm) and incubation at 37°C. Aliquots (2 ml), containing the released drug, were collected at predetermined time intervals (1, 2, 3, 4, 5, 6 and 24 h) and immediately replaced with the same volume of fresh VFS, pH 4.5. The amount of released drug was determined by HPLC.¹⁷

To investigate the influence of pH on AZI release and to mimic cervical conditions, studies were also performed in PBS, pH 7.5. Free AZI dissolved in a mixture of ethanol and demineralized water (6/4, v/v) served as a control and was examined at pH 7.5.

Storage stability studies

The changes in the physical properties of AZI-liposomes (mean diameters, PDIs and zeta potentials) were monitored during storage of the liposome dispersions for 8 months at 4°C. The measurements were performed at 25°C.

Ex vivo vaginal deposition/penetration studies

Porcine vaginal mucosa was used to evaluate the deposition/penetration of AZI from the selected AZI-liposomes. The vaginal tissues from the adult animals (approximately 1 year

old) were received as waste from a local slaughterhouse. The vaginal tissue was carefully separated from the underlying tissue, cleaned with physiological solution (0.9% NaCl), cut into smaller pieces (approximately 3×3 cm) and frozen at –20°C. Approximately 30 min prior to the experiments, the vaginal tissue was defrosted and pre-equilibrated in VFS for 15 min. The vaginal tissue (900–1,050 µm thick) was fixed between the donor and acceptor chamber of the Franz cell (3.14 cm², surface area), with the vaginal epithelium facing the donor chamber. The acceptor chamber (15 ml) was filled with PBS, pH 7.5, and constantly stirred (200 rpm). The whole system was maintained at 37±1°C. Samples of AZI-liposomes or the control (free AZI, 6/4, ethanol/water solution), equivalent to 1 mg of AZI, were added to the donor chamber and covered with parafilm. The acceptor medium (500 µl) was sampled at intervals of 1, 2, 3, 4, 5, 6 and 24 h and immediately replenished with the same volume of fresh PBS. After 24 h, the liquid (if remained) in the donor chamber was removed by pipetting, and the vaginal surface was carefully washed with methanol. Both fractions were mixed together and dissolved in methanol, indicating nonpenetrating AZI. The vaginal tissue was then shred and extracted with methanol (4 h, agitation) to determine the drug that remained inside the vaginal tissue. The AZI content in all the collected samples was quantified by HPLC.¹⁷

Antibacterial assays against *E. coli*

Bacterial strains and growth conditions

The three *E. coli* strains (ATCC 8739, ATCC 700928 and K-12) were stored in 20% glycerol stocks at –70°C. Working stocks of each strain were prepared on LB agar plates and stored at +4°C. For liquid cultures, a 1 µl inoculation loop full of colonies was transferred to 3 ml of LB broth and incubated under aerobic conditions (37°C, 220 rpm) overnight. The overnight cultures were diluted in LB broth and incubated until they reached exponential growth, corresponding to approximately 10⁸ colony forming units per milliliter (CFU/ml). The bacterial concentration was estimated by measuring the turbidity at 595 nm using a Varioskan Flash Multimode plate reader (Thermo Fisher Scientific, Vantaa, Finland) and further confirmed by serially diluting and plating the cultures on LB agar for the determination of CFU/ml.

In vitro antibacterial activity against biofilm and planktonic *E. coli*

The anti-biofilm activity of the selected AZI-liposomes (CL-3, PGL-2 and DPGL-2) toward *E. coli* strains was

tested and compared to the activity of free AZI (6/4, v/v, ethanol/water solution) and the corresponding empty liposomes (without encapsulated AZI).

Two-fold dilution series of AZI-liposomes and free AZI (solution) were prepared in LB and tested at eight concentrations ranging from 0.61 µg/ml to 78.50 µg/ml. The studies were performed in two modes, pre- and post-biofilm formation.²⁴ In both cases, exponentially grown bacterial cultures, diluted to a concentration of 10⁶ CFU/ml, were used. Briefly, in the pre-exposure mode of the assay, bacteria and AZI samples were simultaneously added to NuncTM 96-well polystyrene microplates (NuncTM Delta surface) at a final volume of 200 µl. The plates were incubated at 37°C, 200 rpm, for 18 h. For the postexposure mode, bacteria were initially added to the microtiter well plates (200 µl) and allowed to form biofilms for 18 h (37°C, 200 rpm). After 18 h, the planktonic bacterial phase was discarded, and the AZI samples (prepared as described above) and fresh culture medium were added to the wells. The plates were further incubated for 24 h (37°C, 200 rpm). The samples were tested in duplicate in four independent experiments. Empty liposomes (without AZI) were always included as a negative control, and a mixture of ethanol and water was included as a solvent control, in addition to the wells that contained only medium and untreated bacteria.

Anti-biofilm activity was quantified using a resazurin (redox) staining assay.²⁴ At the end of the incubation periods with the compounds (in the pre-exposure and postexposure modes), the planktonic phase was removed, and the biofilms were washed once with PBS. Resazurin solution in PBS (20 µM) was added to the wells (200 µl), and the plates were incubated at room temperature for 1 h. Fluorescence was measured (λ_{ex} = 560 nm, λ_{em} = 590 nm) using a Varioskan Flash Multimode plate reader using a top fluorescence reading. Anti-biofilm activity was determined as the inhibition percentage compared to untreated biofilms. The concentrations of the samples that inhibited biofilm viability by 50% (IC₅₀) were calculated from eight concentration points via nonlinear regression analysis using GraphPad Prism 6.0 software.

Additionally, antibacterial activity against planktonic bacteria was assessed spectrophotometrically. At the end of the 18 hr incubation period (in the pre-exposure mode of the assay), bacterial turbidity was recorded at 620 nm using a Varioskan Flash Multimode plate reader. Antibacterial effects were determined as inhibition percentages compared to untreated bacteria. The concentrations

of the samples that inhibited bacterial growth by 50% (minimum inhibitory concentrations, MIC₅₀) were determined as described above.

In vitro antibacterial assay against intracellular *C. trachomatis*

Bacterial strain

The *C. trachomatis* serovar D (UW-3/CX, ATCC) strain was propagated and partially purified according to previously described methods.^{25,26} Briefly, DEAE-dextran (45 mg/ml in Hanks' balanced salt solution) treated McCoy cells were infected with *C. trachomatis* serovar D, and incubated for 48 h in Minimal Essential Medium with Earle salts supplemented with 10% fetal bovine serum, 8 mmol/l HEPES, 0.5% glucose, 1× nonessential amino acids, 2 mmol/l L-glutamine, 25 µg/ml gentamicin and 1 µg/ml cycloheximide. Infected cells were lysed by 2 freeze-thaw cycles in sucrose-phosphate-glutamic acid buffer 48 h post-infection. The cell lysate was centrifuged (800 g, 10 min) and the supernatant containing the chlamydial elementary bodies were used for subsequent experiments. The titer of the chlamydial stock was calculated by infecting McCoy cells and counting the inclusion numbers 48 h post-infection with a monoclonal anti-chlamydia LPS antibody (AbD Serotec, Oxford, United Kingdom) and a FITC-labeled anti-mouse IgG (Sigma-Aldrich, St. Louis, MO).

Culture and infection of HeLa cells with *C. trachomatis* in the presence of AZI-liposomes and free AZI

HeLa 229 cells (ATCC) (40,000 cells/well) were cultured in 96-well microtiter plates (Sarstedt, Nümbrecht, Germany) in 100 µl of Minimum Essential Medium supplemented with Earle's salts, 10% heat-inactivated fetal bovine serum, 1× nonessential amino acids, 8 mmol/l HEPES, 2 mmol/l L-glutamine, 1 µg/ml fungisone and 25 µg/ml gentamycin. The cells were incubated overnight at 37°C in 5% CO₂ to obtain a 90% confluent cell layer.²⁷ HeLa cells were washed twice with 100 µl of PBS, pH 7, per well and infected with *C. trachomatis* (multiplicity of infection, MOI 0.2) as described previously.²⁸ After infection, the HeLa cells were washed twice with PBS and cultured in cycloheximide-free medium supplemented with a dilution series of the applied AZI samples (n=3). A concentration range of 0.5–0.0002 µg/ml AZI for AZI-liposomes (CL-3, PGL-2 and DPGL-2) and free AZI (solution) was tested. The cells were incubated for 48 h at 37°C in 5% CO₂.

Cell lysis and direct quantitative polymerase chain reaction (qPCR) measurements

Direct qPCR was performed according to Eszik et al²⁸. Briefly, 48 h post infection, the HeLa cells were washed twice with 100 µl/well PBS. Finally, 100 µl of Milli-Q water (Millipore, Billerica, MA, USA) was added to the wells, and two freeze-thaw cycles were performed. The mixed cell lysates were used as templates in qPCR for chlamydial genomic DNA quantitation. qPCR was performed using the *C. trachomatis* *pykF* gene-specific primers in a Bio-Rad CFX96 real-time system (Bio-Rad, Hercules, CA, USA). The primer sequences were: *pykF* forward: 5'-GTTGCCAACGCCATTACGATGGA-3', *pykF* reverse: 5'-TGCATGTACAGGATGGGCTCCTAA-3'. Two microliters of 5x HOT FIREPol® EvaGreen® qPCR Supermix (Solis BioDyne, Tartu, Estonia), 10 pmol of qPCR primers, 1 µl of cell lysate template and 5 µl of Milli-Q water were used per reaction. The qPCR cycle parameters were as follows: 12 min at 95°C for polymerase activation, 40 cycles at 95°C for 20 s and 64°C for 1 min, and a melting curve analysis step for specificity testing. Cycle threshold (Ct) levels were calculated for each sample. Minimum inhibitory concentration (MIC) values for *C. trachomatis* were identified as previously described.²⁸

In vitro biocompatibility studies

Cell culturing

HeLa cells were cultured as monolayers and maintained in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum, 2 mM L-glutamine, 100 U/ml penicillin and 100 µg/ml streptomycin in a humidified atmosphere with 5% CO₂ at 37°C.

Cell toxicity study

The cells were inoculated onto 96-well microtiter plates at concentrations of 1.6×10^4 cells/ml. AZI-liposomes (CL-3, PGL-2 and DPGL-2) or free AZI (6/4, v/v, ethanol/water solution) was added the next day at five dilutions (39.25 µg/ml, 19.63 µg/ml, 9.81 µg/ml and 2.45 µg/ml and 1.23 µg/ml) and incubated with the cells for an additional 24 h (37°C, 5% CO₂). The cell viability was evaluated by the MTT assay, a colorimetric assay system that measures the reduction of a tetrazolium component (MTT) into an insoluble formazan product by the mitochondria of viable cells.²⁹ The absorbance was determined spectrophotometrically at 570 nm on a microplate reader (Multiscan, Thermo Labsystems, Waltham, MA, USA) and was directly proportional to the cell viability. The

cell viability of the treated cells was expressed as a percentage compared to the untreated control cells. Each result represents an average value from at least three separate experiments performed in quadruplicate.

Statistical analysis

Statistical data analyses were performed using the GraphPad 5.0 Prism program (GraphPad Software, Inc., San Diego, USA). Student's *t*-test was applied for comparisons between two groups, and means were considered significantly different when $p < 0.05$. One-way ANOVA and Tukey's multiple comparison test, with $p < 0.05$ set as the minimal level of significance, were applied for comparisons of three or more groups.

Results and discussion

Characterization of liposomes

A suitable characterization of liposomes is crucial for obtaining reliable data with high translational output.³⁰ Namely, the physicochemical properties of liposomes influence their stability in biological environments and during storage, determine the pharmacokinetic properties of the encapsulated drug and contribute toward their behavior within the biological milieu, subsequently affecting the efficiency of therapy. Therefore, the first phase in the present study focused on assessing the physical characteristics of AZI-liposomes that differed in terms of bilayer composition, the presence of cosolvent and the edge activator (Table 1). Liposomes were evaluated for size and surface charge (Tables 2 and 3), bilayer elasticity (Table 4) and encapsulation efficiency (Figure 1).

As presented in Table 2, AZI-liposomes composed of only neutral phospholipids (CL-2, PGL-1 and DPGL-1) were significantly larger (544–840 nm), with broad size distributions ($PDI > 0.49$), than the corresponding liposomes containing negatively charged EPG (CL-3, PGL-2 and DPGL-2) (*t*-test, $p < 0.05$). Extrusion of all AZI-liposomes significantly decreased their mean diameters (189–261 nm) and PDIs (0.17–0.27) (*t*-test, $p < 0.05$). The only exception was PGL-1 with $PDI > 0.7$, indicating a tendency of PGL-1 to form aggregates in the presence of propylene glycol and the absence of a negatively charged phospholipid, which both affect the size of the liposomes. Comparison of the various types of negatively charged liposomes (CL-1, CL-3, PGL-2 and DPGL-2) revealed DPGL-2 as the smallest. These results are in agreement with previously published findings showing the ability of

Table 2 Size and surface characteristics of the AZI-liposomes

Liposomes	Originally prepared		Extruded (3x400 nm)		Zeta potential (mV)
	Mean diameter (nm)	PDI	Mean diameter (nm)	PDI	
CL-1	351±3	0.27±0.02	259±1*	0.17±0.01*	-66.4±1.3
CL-2	840±11	0.63±0.08	221±2*	0.26±0.10*	-21.6±0.6
CL-3	466±3	0.24±0.03	262±3*	0.18±0.03	-61.5±0.7
PGL-1	545±14	0.56±0.01	341±27 [#] *	0.74±0.08*	-18.2±0.5
PGL-2	529±10	0.24±0.01	222±2*	0.20±0.02*	-55.8±0.9
DPGL-1	574±8	0.49±0.09	219±1*	0.27±0.01*	-17.8±0.6
DPGL-2	287±3	0.31±0.01	189±1*	0.22±0.02*	-53.1±0.5

Notes: The values are the mean ± S.D.; n=3 (mean diameter, PDI); n=5 (zeta potential); [#]Estimated size due to high PDI value (>0.7). *Statistically significant compared to the originally prepared liposomes (t-test, $p<0.05$).

Abbreviation: PDI, polydispersity index.

Table 3 Characteristics of the AZI-liposomes at simulated vaginal conditions

Liposomes	Mean diameter (nm)	PDI	Zeta potential (mV)
CL-1	276±5 ^a 270±3 ^b	0.19±0.02 ^a 0.20±0.02 ^b	-0.6±0.3 ^a -0.4±0.1 ^b
CL-2	392±24 ^a 397±12 ^b	0.46±0.08 ^a 0.42±0.10 ^b	4.1±0.4 ^a 4.4±0.3 ^b
CL-3	323±4 ^a 331±3 ^b	0.22±0.02 ^a 0.21±0.09 ^b	-4.4±0.3 ^a -4.2±0.3 ^b
PGL-1	382±7 ^a 389±9 ^b	0.77±0.09 ^a 0.73±0.08 ^b	4.4±0.3 ^a 5.1±0.4 ^b
PGL-2	274±4 ^a 276±3 ^b	0.15±0.03 ^a 0.17±0.08 ^b	-1.8±0.3 ^a -2.1±0.1 ^b
DPGL-1	262±5 ^a 258±3 ^b	0.21±0.10 ^a 0.22±0.05 ^b	3.8±0.3 ^a 4.6±0.2 ^b
DPGL-2	217±2 ^a 212±3 ^b	0.19±0.05 ^a 0.20±0.03 ^b	-1.6±0.2 ^a -2.9±0.2 ^b

Notes: ^a37°C; ^b25°C. The extruded AZI-liposomes were dispersed in VFS, pH 4.5, and the mean diameters, PDIs and zeta potentials were determined at 37°C. The measurements were also performed at 25°C to investigate the effect of temperature on the physical properties of liposomes. The each value represents the mean ± S.D. (n=3, mean diameter, PDI; n=5, zeta potential). Significant differences between the determined parameters were not obtained when experiments were carried out at 25 or 37°C (t-test, $p>0.05$).

Abbreviation: PDI, polydispersity index.

an edge activator to reduce the size of deformable liposomes,^{21,23} which might be attributed to the enhanced elasticity (flexibility) and decreased surface tension of the liposomes in the presence of single chain surfactants.³¹ Moreover, a decrease in the size of microemulsions containing an increased ratio of monoacyl phosphatidylcholine (used as the edge activator in DPGLs in the present study) has been confirmed by Hoppel et al³².

The zeta potential values (Table 2) followed the properties of the materials used for the preparation of liposomes. Liposomes prepared without EPG had zeta potentials between -17 and -21 mV, while those containing EPG

had strongly negative zeta potentials higher than -50 mV, indicating the formation of physically stable liposome dispersions.

To resemble more closely the conditions AZI-liposomes will be exposed to, the evaluation of their physicochemical properties was also performed at simulated vaginal conditions (VFS, pH 4.5, 37°C). Moreover, to determine the possible effect of the temperature on the size and surface charge of AZI-liposomes, the experiments were also conducted at 25°C.

The results presented in Table 3 demonstrate significant increase (t-test, $p<0.05$) in mean diameters of all AZI-liposomes because of their interaction with vaginal fluid

Table 4 Liposome bilayer elasticity

Liposomes	J (g)	r_v (nm)	E
CL-1	1.61±0.2	165±2	4.4±0.3*
CL-2	0.17±0.0	185±3	0.58±0.1
CL-3	0.19±0.1	165±2	0.52±0.2
PGL-1	4.26±0.3	178±1	13.53±0.3*
PGL-2	4.41±0.2	169±2	12.64±0.2*
DPGL-1	6.63±0.4	154±2	15.70±0.4*
DPGL-2	7.91±0.3	147±1	17.02±0.3*

Notes: The values are the mean ± S.D. (n=3). One-way ANOVA and Tukey's multiple comparison test, with $p < 0.05$ set as the minimal level of significance, were applied for comparison of the E values. *Statistically significant compared to CL-2 and CL-3 ($p < 0.05$).

Abbreviations: E, degree of liposome bilayers elasticity; J, amount of extruded liposomes; r_v , average diameter of liposomes following extrusion under constant pressure.

components, in comparison to the corresponding values shown in Table 2. However, polydispersity indexes commonly remained unchanged despite of the used media (buffer, pH 7.5 or VFS, pH 4.5), except for the CL-2 and PGL-1.

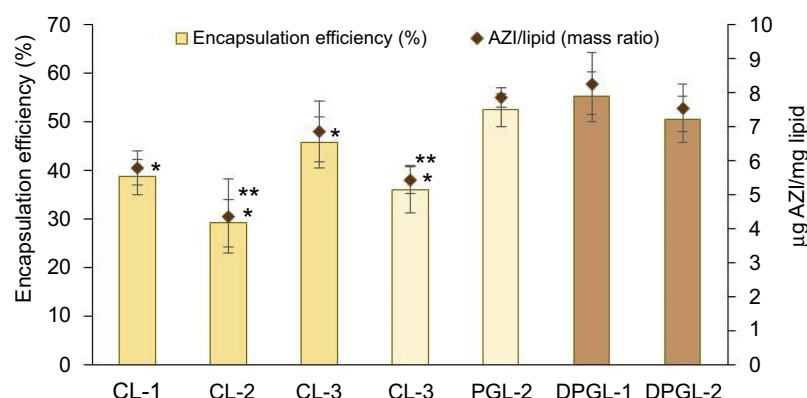
Similarly as the mean diameters, the zeta potentials of AZI-liposomes (Table 2) have significantly changed (t -test, $p < 0.05$) at mimicked vaginal conditions (Table 3). The originally, highly negative zeta potentials of CL-1, CL-3, PGL-2 and DPGL-2 decreased from approximately -50 mV (buffer, pH 7.5) to -4 mV (VFS, pH 4.5), while AZI-liposomes with slightly negative surface charge (between -18 and -21 mV) changed to slightly positive values (4 mV). Interestingly, the temperature did not affect the size and surface properties of the liposomes (t -test, $p > 0.05$).

Since the cervical tissue is characterized by neutral pH³³ and considering the fact that the temperature had no impact on the physical properties of liposomes (Table 3), the physical properties of AZI-liposomes shown in Table 2 (obtained

in the buffer, pH 7.5) could be translated to mimicked cervical conditions.

The thermodynamic state of the liposome bilayers is responsible for their release pattern and permeation ability, as well as the interactions of liposomes with bacteria and biofilms.^{17,23,34} Liposome bilayer elasticity/rigidity was evaluated by applying an easy and reproducible method based on the passage of liposomes through 100-nm membranes under a constant external pressure (2.5 bar) for 5 min.²² The results presented in Table 4 confirmed that PGLs and DPGLs are elastic liposomes, while CLs, particularly CL-2 and CL-3, exhibited pronounced rigid bilayers. In addition, the degree of bilayer elasticity (E) was confirmed to be commonly dependent on the quantity of extruded liposomes (J) rather than their size following extrusion (r_v) through 100-nm membranes (r_p). Comparison of the calculated E values among the different CLs showed the lowest elasticity for CL-2 and CL-3 due to the presence of 25% (w/w) hydrogenated soybean phosphatidylcholine (SPC-3) in the bilayers. In contrast, PGLs and DPGLs were characterized by flexible bilayers with E values between 12 and 17, whereas the elasticity of PGLs was somewhat less pronounced. The highest E value was confirmed for DPGL-2 (ANOVA, $p < 0.05$), with a value almost 34-fold higher than the E values of CL-2 and CL-3. The increased elasticity of DPGLs compared with PGLs is ascribed to the presence of 15% (w/w) monoacyl phosphatidylcholine in DPGLs. These results are in accordance with previously published findings for DPGLs containing the same ratio of propylene glycol but with sodium deoxycholate as an edge activator.²²

Properly balancing physical characteristics enables the development of liposomes with suitable encapsulation of AZI as a prerequisite for adequate therapeutic outcomes.

**Figure 1** Encapsulation of AZI in different types of liposomes expressed as the encapsulation efficiency (%) and the drug-per-lipid ratio.

Notes: The values indicate the mean ± S.D. *Significantly different from DPGL-1 (ANOVA, $p < 0.05$). **Significantly different from the corresponding highly negatively surface-charged liposomes (t -test, $p < 0.05$).

Abbreviation: AZI, azithromycin.

Since our aim was topical vaginal therapy, liposomes with diameters of 200–300 nm are expected to be optimal.³⁵

As shown in Figure 1, encapsulation of AZI differed between 29% (4.4 µg AZI/mg lipid) for CL-2 and 55% (8.3 µg AZI/mg lipid) for DPGL-1. These results are a consequence of the physicochemical properties of AZI and the liposomal constituents, both affecting encapsulation of the drug. AZI, even in the form of dihydrate, exhibits lipophilic features (log $P=3.98$),¹³ therefore it is expected to become accommodated within the liposome bilayers. However, molecular weight of AZI is rather large (Mw 785),¹³ restricting its incorporation within bilayers. The differences in the encapsulation efficiencies observed between the various types of liposomes thus proceed from the effects of the liposomal constituents on AZI incorporation. Generally, liposomes with EPG embedded as a bilayer constituent were able to incorporate significantly larger amounts of the drug (except for DPGL-2) than the corresponding liposomes composed of only neutral phospholipids (*t*-test, $p<0.05$). Such results are probably a consequence of the interaction between AZI and the negatively charged EPG inside liposome bilayers, facilitating the encapsulation of AZI. Binding of AZI to negatively charged phospholipids has been confirmed by Van Bambeke et al³⁶. Among the various negatively surface-charged liposomes (CL-1, CL-3, PGL-2 and DPGL-2), both types of elastic liposomes (PGL-2 and DPGL-2) demonstrated significantly higher encapsulation efficiency than CL-1 and CL-3 (ANOVA, $p<0.05$). Increased incorporation of AZI in the elastic liposomes was due to the presence of propylene glycol (PGLs, DPGLs) and the monoacyl phospholipid SLPC-80 (DPGLs), which both facilitate solubilization and hence improve encapsulation of AZI. Contrary to the previous findings obtained for deformable liposomes,^{17,21,23} DPGLs exhibited superior drug encapsulation because the presence of monoacyl phosphatidylcholine increased the solubilization of AZI. Namely, monoacyl phospholipid has been proven as a more efficient solubilizer than diacyl phospholipid.³⁷ The slightly lower encapsulation of AZI in DPGL-2 than in DPGL-1 could be attributed to the possible interaction/competition of the monoacyl phospholipid and the negatively charged phospholipid inside the bilayers of DPGL-2 for AZI; however, further studies, such as Fourier transform infrared spectroscopy (FT-IR), are necessary to prove the interactions of liposomal bilayer constituents and the drug.

In vitro release studies

The predictable release of the encapsulated drug from liposomes in medium simulating the in vivo environment

is of great significance regarding the use of liposomes as drug delivery nanosystems. As the human vaginal mucosa has a pH between pH 4 and 5, studies were performed in conditions simulating the vaginal environment, ie, VFS, pH 4.5. The experiments were also performed in PBS, pH 7.5, to mimic cervical conditions,³³ and to investigate the effect of pH and vaginal fluid components on AZI release from the liposomes.

The results in Figure 2 demonstrate the prolonged release of AZI from all the liposomes in both tested media relative to the control (AZI solution) (ANOVA, $p<0.05$). Overall, the release from all the tested liposomes was slightly increased when the liposomes were dispersed in VFS, pH 4.5, than in PBS, pH 7.5, as a result of the effects of lower pH and vaginal fluid components on liposome stability.³⁸ Despite the used release media, all the liposomes exhibited initial burst release followed by sustained AZI release. Such a release profile might be relevant for facilitating the rapid start of antibiotic activity and ensuring increased local drug concentrations.

Liposome membrane elasticity/rigidity significantly influenced the AZI release profile from the different liposomes under simulated cervicovaginal conditions. Thus, the slowest release, at both pH 4.5 and 7.5, was obtained from the rigid liposomes, ie, CL-2 and CL-3, followed by less rigid CL-1, while AZI release was significantly faster from both types of elastic liposomes. For instance, 70–80% of the originally encapsulated AZI was released after 6 h from PGLs and DPGLs under simulated vaginal conditions, compared to 40–45% with CL-2 and CL-3. Increased AZI release from PGLs, especially from DPGLs, was expected due to the increased membrane permeability caused by propylene glycol (PGLs) in addition to intercalation of the edge activator (SLPC-80) in bilayers of DPGLs. These observations are in accordance with previous reports for PGLs and DPGLs.^{17,22,39} In addition to bilayer elasticity, the highly negative surface charge also increased the release of AZI. Similar behavior of negatively charged liposomes was also observed by Pavelić et al³⁸. However, this effect was negligible in comparison to the impact of membrane elasticity (Figure 2).

Storage stability studies

The stability of liposomes during storage is an important feature considering their use as drug nanoformulations. As shown in Figure 3, all the liposomes with slightly negative surface charges (CL-2, PGL-1, and DPGL-1) significantly

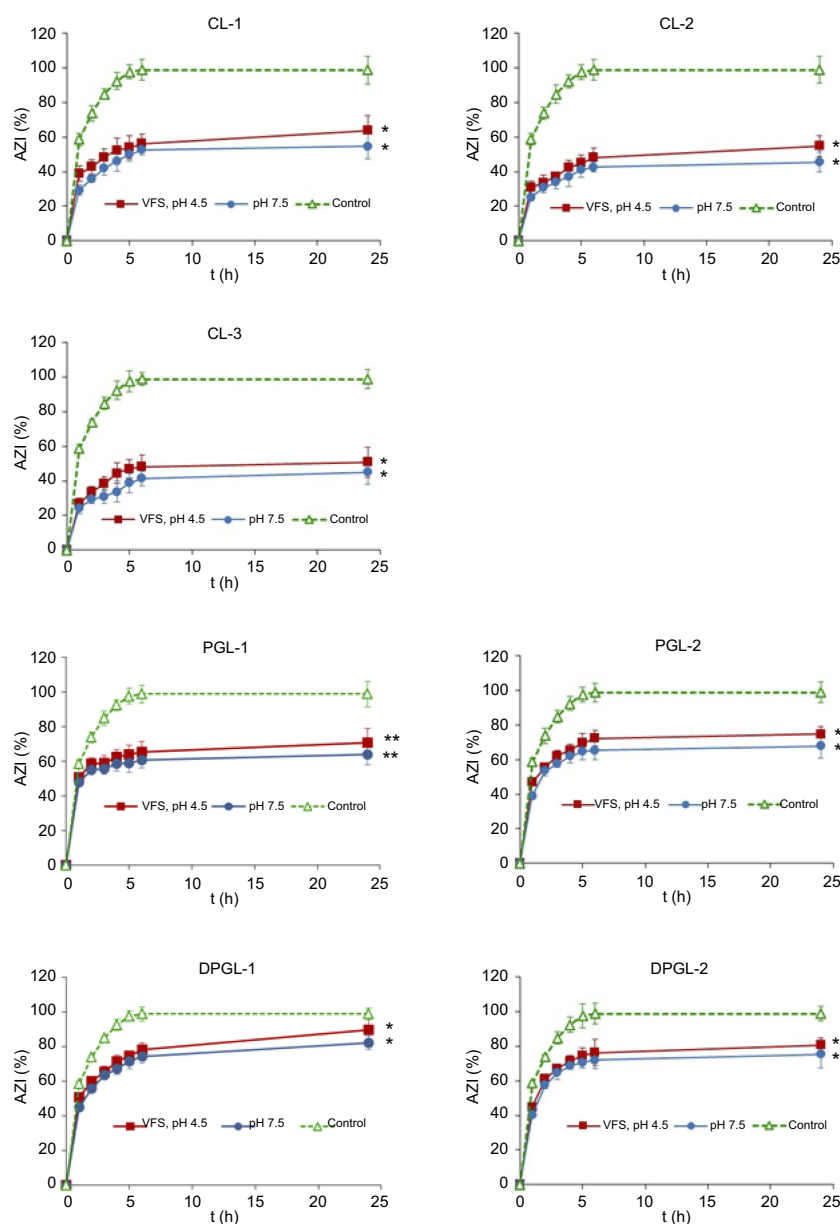


Figure 2 In vitro AZI release from the different types of liposomes.

Notes: The each value represents the mean \pm S.D. ($n=3$). One-way ANOVA and Tukey's multiple comparison test, with $p<0.05$ set as the minimal level of significance, were applied for comparison of AZI release from the different types of liposomes and control. The control, representing a solution of AZI, was examined at pH 7.5. *Significantly different (ANOVA, $p<0.05$) from the control (period 1–24 h). **Significantly different (ANOVA, $p<0.05$) from the control (period 2–24 h).

Abbreviation: AZI, azithromycin.

increased in size (mean diameters and PDIs) during storage in comparison to the corresponding initial values; CL-2 and DPGL-1 increased after 2 months (CL-2 from 221 to 267 nm and DPGL-1 from 248 to 274 nm), while PGL-1 increased after 5 months (from 341 to 372 nm) (t -test, $p<0.05$) (Figure 3A and B). The lower physical stabilities of CL-2 and DPGL-1 were closely associated with their slightly negative surface charges (-20 and -18 mV, Figure 3C). In addition, DPGL-1 was characterized by the significantly pronounced membrane elasticity (15.7, Table 3), making their bilayers

more permeable and prone to aggregation (fusion). These observations are consistent with the recently reported findings demonstrating the influence of the bilayer elasticity on the reduced physical stability of liposomes incorporating lipophilic drug.¹⁷ In contrast, the liposomes with highly negative surface charge (zeta potentials between -50 and -60 mV), namely, CL-1, CL-3, PGL-2 and DPGL-2, were physically stable during 7 months of storage (Figure 3C). Similar findings were previously reported by Palac et al,²¹ whereas highly negatively surface charged elastic and rigid

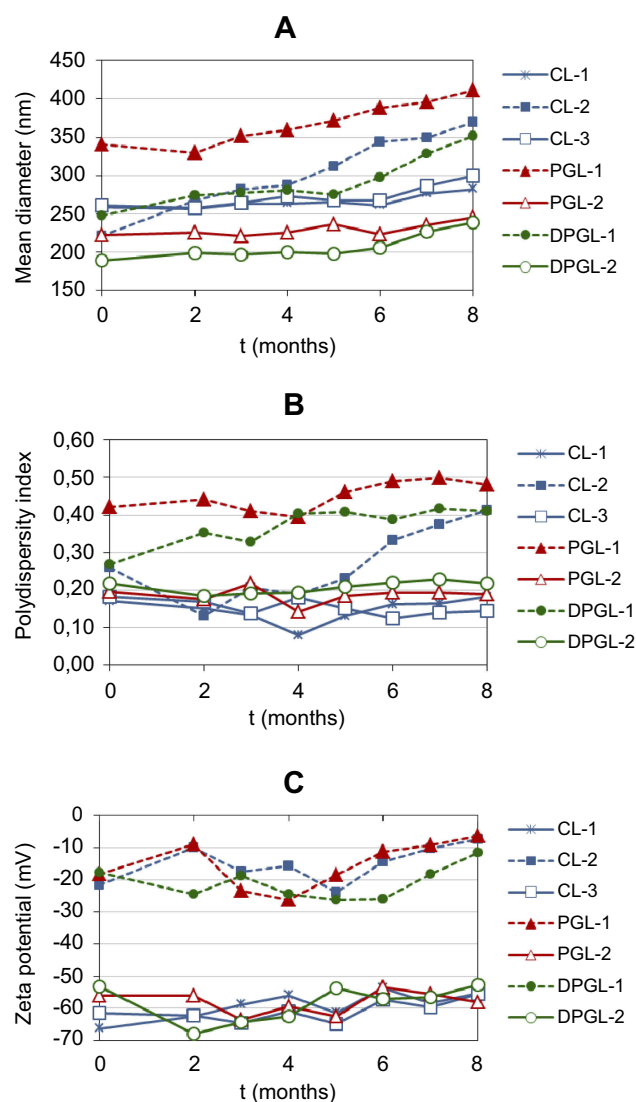


Figure 3 Physical stability of different AZI-liposomes: mean diameters (A), polydispersity indexes (B) and zeta potentials (C).

Notes: The each value represents the mean \pm S.D.; $n=3$ (mean diameter, polydispersity index); $n=5$ (zeta potential). For each type of AZI-liposomes, the values determined during 2–8 months were compared with the corresponding initial values using Student's *t*-test, with $p < 0.05$ set as the minimal level of significance. The standard deviations and statistically significant differences are not shown due to the clarity of presentation. Statistically significant increase in mean diameters and polydispersity indexes are determined for CL-2 and DPGL-1 (2–8 months) and for PGL-1 (5–8 months).

liposomes with encapsulated hydrophilic drug remained physically stable over 10 months of storage at 4°C.

A comparison of the two different preparations of highly negatively charged CLs (CL-1 and CL-3) showed similar stability profiles. Since the bilayers of CL-3 were significantly more rigid than those of CL-1 (Table 3), it is assumed that CL-3 will be more suitable for further studies due to the lower possibility of drug leakage.

Finally, taking into account the physical characteristics of the liposomes, suitable AZI encapsulation, in vitro AZI release

patterns under simulated cervicovaginal conditions and storage stability, CL-3, PGL-2 and DPGL-2 were selected for further investigations. Their composition is schematically depicted by Figure 4. Besides phosphatidylcholine (major phospholipid) and 5% phosphatidylglycerol (negatively charged lipid) common for all optimized liposomes (CL-3, PGL-2 and DPGL-2), the bilayers of CL-3 were enriched with hydrogenated phosphatidylcholine (SPC-3) contributing to membrane rigidity. On the other hand, elastic liposomes embedded either propylene glycol (PGL-2) or both propylene glycol and monoacyl phospholipid (DPGL-2) in their bilayers, decreasing bilayers' compactness, while increasing its elasticity.

Ex vivo permeation studies

Localized antimicrobial therapy has been used to treat vaginal infections. However, its efficiency is frequently reduced because of several factors. For instance, possible systemic drug absorption and drug teratogenicity (metronidazole) can take place, and inadequate local drug concentrations may be achieved, along with the possible development of antibiotic-resistant strains, the formation of biofilms and the failure of the drugs to reach the desired site of action. These possibilities make oral administration of high doses of antibiotics necessary, which may lead to an increased incidence of undesired drug side effects, patient inconvenience and an enhanced risk of antibiotic resistance development.⁴⁰

The use of liposomes represents a suitable approach to achieve effective localized vaginal therapy. Manipulation of the physical characteristics of liposomes (bilayer elasticity/rigidity, surface coating) enables the delivery of encapsulated drugs to the site of action.¹⁰ Therefore, to more closely resemble the applicability of AZI-liposomes for the treatment of vaginal infections, *ex vivo* permeation studies were performed.

The results presented in Figure 5 refer to AZI levels determined in the acceptor chamber (penetrated AZI), within vaginal tissue, and on the vaginal surface (nonpenetrated AZI) after 24 h of testing. It was not possible to detect the presence of AZI in the acceptor chamber during the first 6 h of testing since the drug content was under the limit of detection. The only exception was the control (free AZI), for which a negligible amount of the drug was detected after 5 h.

The performed *ex vivo* studies undoubtedly demonstrate the potential of AZI-liposomes in localized vaginal therapy (Figure 5). All tested liposomes enabled accumulation of the drug on the vaginal surface (17–24%) and mostly inside

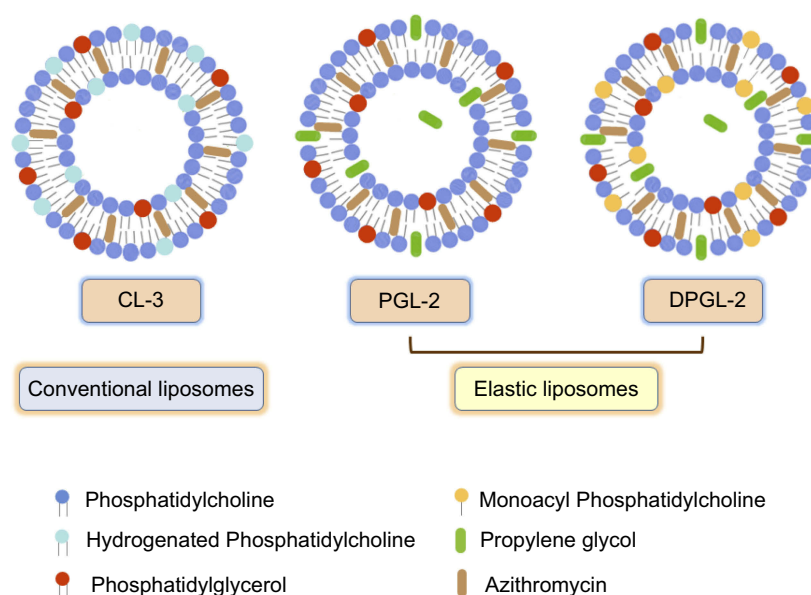


Figure 4 Schematic drawing of the AZI-liposomes explored for ex vivo vaginal deposition/penetration, in vitro antibacterial activity and biocompatibility studies.

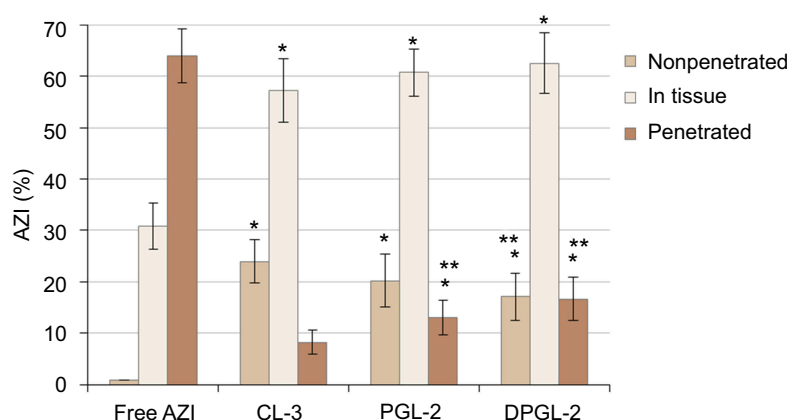


Figure 5 Ex vivo deposition/penetration of AZI into/through porcine vaginal tissue.

Notes: The each value represents the mean \pm S.D. ($n=3$). One-way ANOVA and Tukey's multiple comparison test, with $p<0.05$ set as the minimal level of significance, were applied for comparison of the levels of nonpenetrated, penetrated and AZI retained in the tissue, from different types of AZI-liposomes and control. *Significantly different (ANOVA, $p<0.05$) from free AZI (control). **Significantly different (ANOVA, $p<0.05$) from CL-3.

Abbreviation: AZI, azithromycin.

vaginal tissue (57–63%), generating higher local concentrations of AZI and reducing its undesired penetration to the systemic circulation compared to the control (free AZI). Increased retention of the drug inside the vagina could be relevant for the treatment of recurrent bacterial vaginosis and biofilm-related infections. Comparison of the different types of liposomes demonstrated the highest penetration ability for DPGL-2, followed by PGL-2 and CL-3 (Figure 5). These findings are consistent with the bilayer properties of liposomes (Table 3) influencing AZI deposition/penetration in vaginal tissue. For instance, CL-3 had rigid bilayers facilitating retention of the AZI on/within the vaginal tissue. In

contrast, the highest AZI penetration was observed with DPGL-2 (Figure 5), which is in accordance with its very elastic bilayers, facilitating increased drug penetration into/through the vaginal tissue.

The physicochemical properties of liposomes significantly contribute to the vaginal deposition/penetration abilities of the encapsulated drug. Thus, mucoadhesive, chitosan-coated liposomes enabled the localization of clotrimazole on the vaginal surface,⁴¹ while non-mucoadhesive, ie, mucus-penetrating, PEGylated liposomes demonstrated increased interferon alpha penetration into/through the vaginal mucosa,⁴² in comparison to controls (solutions of the free drugs at the same

concentrations present in the tested liposomes). In a recent study, positively charged liposomes with rigid bilayers enabled localization of AZI on the skin surface due to strong interactions with the negatively charged skin cells.¹⁷

In vitro antibacterial activity

The optimized AZI-liposomes (CL-3, PGL-2 and DPGL-2) were investigated for their in vitro antibacterial activity against several biofilm-forming and planktonic *E. coli* strains (Tables 5 and 6) and against intracellular *C. trachomatis* (Figure 6).

The results for planktonic *E. coli* are expressed as MIC₅₀ values, while those obtained for the pre-exposure mode of the biofilm assay are presented as IC₅₀ values (Table 5).

As shown in Table 5, the antibacterial activity of all the AZI samples (both liposomally encapsulated and free AZI) varied among the strains. In general, *E. coli* K-12 was the most sensitive to AZI, followed by *E. coli* ATCC 700928 and ATCC 8739. All the AZI-liposomes were more effective than the free AZI against *E. coli* ATCC 700928, with MIC₅₀ values approximately 3-fold lower than the MIC₅₀ of the free AZI. A comparison of the activities achieved by the different types of liposomes toward *E. coli* K-12 demonstrated the similar efficacy of CL-3 and PGL-2, while the activity of DPGL-2 was significantly lower (ANOVA, $p < 0.05$). Regarding *E. coli* ATCC 8739, all of the AZI-liposomes demonstrated activity similar to that of the free antibiotic; CL-3 exhibited slightly lower activity, but this activity was not significantly different from that of free AZI (t -test; $p > 0.05$).

Similarly, the observed anti-biofilm activity of the AZI-liposomes varied between strains. For example, CL-3 was the most effective against *E. coli* ATCC 700928 and K-12, exhibiting IC₅₀ values almost 8-fold lower than those of the free antibiotic. PGL-2 and DPGL-2 demonstrated lower anti-biofilm activity than CL-3 (ANOVA, $p < 0.05$), but this activity was still significantly stronger than that of the free AZI (t -test, $p < 0.05$). PGL-2 inhibited *E. coli* ATCC 8739 biofilm formation with an IC₅₀ of 6.36 µg/ml, which was twice as low as the IC₅₀ of the free AZI (Table 5). By contrast, CL-3 and DPGL-2 displayed weaker anti-biofilm activity (Table 5).

The observed variability in the activity of the different types of AZI-liposomes tested might be a consequence of the bilayer properties of the tested liposomes affecting AZI release and the susceptibility of the different *E. coli* strains to AZI. CL-3 contained hydrogenated phospholipids in the bilayers, contributing to a higher phase transition temperature,⁴³

Table 5 Antimicrobial activity of AZI-liposomes against planktonic bacteria and biofilms in the pre-exposure mode of the assay, as measured by turbidity measurements and resazurin staining, respectively

Sample	MIC ₅₀ (µg/ml) (95% confidence intervals)				IC ₅₀ (µg/ml) (95% confidence intervals)			
	Planktonic bacteria				Prevention of biofilm formation			
	<i>E. coli</i> ATCC 8739	<i>E. coli</i> ATCC 700928	<i>E. coli</i> K-12		<i>E. coli</i> ATCC 8739	<i>E. coli</i> ATCC 700928	<i>E. coli</i> K-12	
Free AZI	7.50 (5.89–9.56)	7.51 (6.66–8.47)	2.76 (1.57–4.82)		14.08 (12.76–21.05)	21.09 (19.16–23.29)	10.01 (8.79–12.21)	
CL-3	8.42 (5.90–9.56)	2.48 (1.95–3.14)	1.66 (1.54–2.31)		45.65 (40.23–55.27)	2.70 (2.11–4.04)	1.26 (1.13–1.85)	
PGL-2	7.38 (4.62–11.81)	2.31 (1.62–3.29)	1.26 (0.66–1.74)		6.36 (5.46–10.46)	12.15 (11.03–14.22)	3.49 (1.87–6.55)	
DPGL-2	7.66 (6.97–10.79)	2.49 (1.69–3.68)	2.61 (1.59–4.27)		26.65 (24.56–36.51)	11.24 (7.79–16.23)	6.10 (4.47–8.32)	

Abbreviations: IC₅₀, half-maximal inhibitory concentration; MIC₅₀, concentration of the antibiotic that inhibited bacterial growth by 50%.

Table 6 Eradication of *E. coli* biofilms at the highest tested concentration (78.50 µg/ml) in the postexposure mode of the assay, as measured using a resazurin staining assay

Sample	Inhibition of biofilm viability (%)		
	<i>E. coli</i> ATCC 8739	<i>E. coli</i> ATCC 700928	<i>E. coli</i> K-12
Free AZI	17.49±1.54	26.44±7.14	26.88±6.00
CL-3	16.44±2.31	25.15±4.83	26.95±3.06
PGL-2	26.94±1.86	27.15±5.09	30.31±1.55
DPGL-2	25.24±14.01	45.67±7.14*	38.08±13.74**

Notes: The values indicate the mean ± S.D. (n=4). *Statistically significant compared to free AZI, PGL-2 and CL-3 (ANOVA, $p<0.05$). **Statistically significant compared to free AZI, PGL-2 and CL-3 (ANOVA, $p<0.05$).

Abbreviation: AZI, azithromycin.

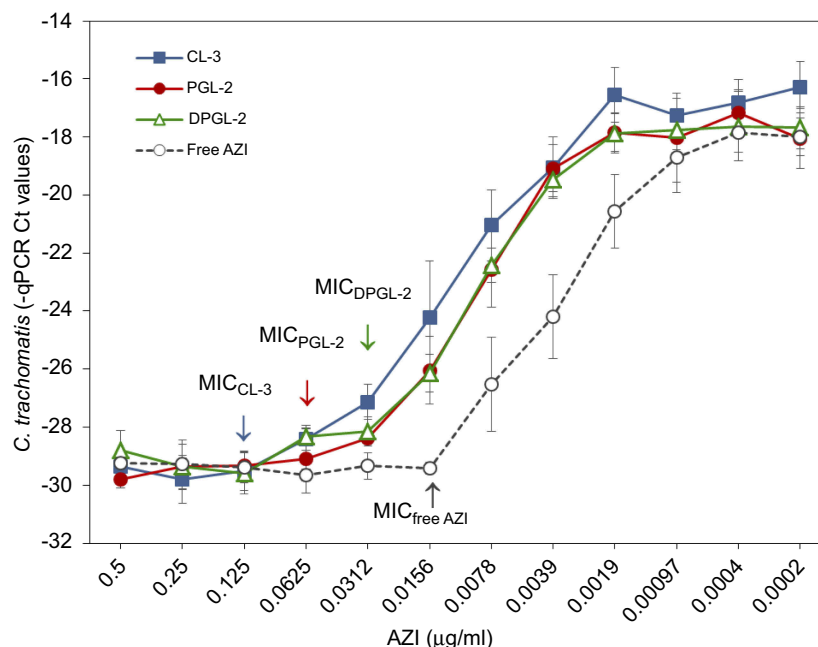


Figure 6 In vitro activities of AZI-liposomes against *C. trachomatis* presented as MIC values.

Notes: *C. trachomatis* (MOI 0.2)-infected HeLa cells were cultured in the presence of 0.5–0.0002 µg/ml free or liposomally loaded AZI (CL-3, PGL-2 and DPGL-2). *C. trachomatis* genome concentrations were measured at 48 h post infection by direct qPCR. Data are presented as the mean Ct levels ± S.D. (n=3).

Abbreviations: AZI, azithromycin; Ct, cycle threshold; MIC, minimum inhibitory concentration; qPCR, quantitative polymerase chain reaction.

allowing slower AZI release, which could be relevant for preventing biofilm formation by *E. coli* ATCC 700928 and K-12.

To examine the potential of AZI-liposomes to eradicate preformed *E. coli* biofilms, the biofilms were exposed to AZI samples (both liposomally encapsulated and the free drug).

As shown in Table 6, the strongest inhibition of the *E. coli* biofilms was achieved with DPGL-2, even though this formulation showed lower activities than CL-3 and PGL-2 against planktonic *E. coli* and toward the suppression of biofilm formation (pre-exposure mode). DPGL-2 inhibited *E. coli* ATCC 700928 biofilm viability by 46% compared with the 27% and 26% inhibition obtained with PGL-2 and the free AZI, respectively. However, its anti-biofilm activity

varied between strains. The strongest inhibition (ANOVA, $p<0.05$) was obtained against *E. coli* ATCC 700928 biofilms, followed by *E. coli* K-12 and *E. coli* ATCC 8739 biofilms (Table 6). When tested against *E. coli* ATCC 8739, PGL-2 was the most active among the different AZI-liposomes (27% inhibition), but it was not significantly more active versus DPGL-2 (25% inhibition, ANOVA, $p>0.05$). However, PGL-2 displayed significantly enhanced activity compared to the free antibiotic (17%), while the anti-biofilm activity of CL-3 was similar to that of the free drug (Table 6).

Overall, anti-biofilm activity (Table 6) was lower than in the pre-exposure mode of the assay (Table 5). These findings are in agreement with the well-known fact that eradication of bacterial biofilms is difficult to achieve, even with

the application of very high concentrations of antibiotics.⁴⁴ The enhanced activities of elastic liposomes, particularly DPGL-2 against *E. coli* biofilms, were probably a consequence of the bilayer properties of the elastic liposomes, which permit their enhanced penetration into the formed biofilms. Additionally, the presence of monoacyl phosphatidylcholine in DPGL-2 has been shown to increase antibiofilm activity.

A limited number of studies reported on the efficacy of liposomal antibiotics against planktonic *E. coli*, but none examined activity against *E. coli* biofilms. According to a literature search, only so-called fusogenic liposomes encapsulating tobramycin or vancomycin have been examined against planktonic *E. coli*.^{45,46} Their superior activities in comparison to the free antibiotics were considered a result of the confirmed fusion of the vesicles composed of dioleoyl phosphatidylethanolamine, dipalmitoyl phosphatidylcholine and cholesteryl hemisuccinate (4:2:4, molar ratio) with *E. coli*, allowing release of the encapsulated vancomycin in the cytoplasm of *E. coli*.⁴⁶

The obtained data for in vitro antibacterial activity (Tables 5 and 6) are important regarding the potential use of AZI-liposomes in localized vaginal therapy for *E. coli* infections. CL-3 and PGL-2 would have potential for the treatment of superficial infections caused by planktonic *E. coli* and the prevention of complicated infections, while DPGL-2 seems the most promising among the tested liposomes for the treatment of biofilm-related infections. The AZI-liposomes with elastic bilayers exhibited enhanced antibacterial activities against *Staphylococcus aureus* and MRSA clinical isolates,¹⁷ which is of great relevance considering the possible use of AZI-liposomes for the treatment of aerobic vaginitis caused by both *E. coli* and *S. aureus*. Cervicovaginal infections caused by *S. aureus* have also been reported to increase the risk of fetal complications in pregnant women, similar to *E. coli*.⁴⁷ In the present study, all AZI-liposomes were negatively charged, which could be significant regarding the biocompatibility issue. Interestingly, the negative surface charge has also been shown to influence the antimicrobial efficacy of AZI-liposomes toward *M. avium*, as demonstrated by Oh et al.¹⁶ Since AZI is a potent broad-spectrum antibiotic, we assume that AZI-liposomes could be suitable for the treatment of recurrent bacterial vaginosis caused by bacteria other than *E. coli* and *S. aureus*, based on the reported efficacy of AZI against *Gardnerella vaginalis*, *Atopobium vaginae*, *Bacteroides* spp., *Mobiluncus* spp. and *Mycoplasma* spp.^{48,49} Moreover, high doses of AZI are

recommended as the first choice of therapy for the treatment of *A. vaginae* and *G. vaginalis* bacterial vaginosis to prevent preterm birth.⁵⁰ Therefore, it seems reasonable to broaden the possible applicability of AZI-liposomes for the treatment of recurrent bacterial vaginosis; however, the efficacy of the developed liposomes has to be investigated to prove these assumptions.

In addition to *E. coli*, the AZI-liposomes were also tested against intracellular *C. trachomatis*. A direct qPCR method was applied to determine the MICs of the free AZI and the liposomally loaded AZI (CL-3, PGL-2, and DPGL-2). *C. trachomatis* (MOI 0.2)-infected HeLa cells were cultured in the presence of 0.5–0.0002 µg/ml AZI samples, and the dose-response curves were evaluated 48 h post infection (Figure 6). The MIC values were identified by statistical comparison of the bacterial DNA concentrations (Ct values) measured in the presence of a given AZI concentration with the Ct values of bacterial DNA in the presence of the highest AZI concentration. The Ct levels measured in the presence of the highest antibiotic concentrations were considered as the genome concentration of the inoculum. The MIC value was defined as the lowest AZI concentration, where the Ct values were not significantly different from the inoculum.²⁸

Among the different types of AZI-liposomes tested against *C. trachomatis*, DPGL-2 was the most effective, with MIC value of 0.0312 µg/ml, followed by PGL-2 (0.0625 µg/ml) and CL-3 (0.125 µg/ml) (Figure 6). These results correlate with the bilayer elasticity/rigidity of the examined liposomes (Table 4), leading to more rapid AZI release from DPGL-2 than CL-3 (Figure 2). The lower efficacy of all the AZI-liposomes than the control (free AZI) could be a consequence of the fact that *C. trachomatis* is an intracellular pathogen and that lipophilic AZI in solution form is more easily taken up by the cells than liposomes,^{51,52} which was also confirmed by *ex vivo* penetration studies (Figure 5), showing higher penetration of free AZI through the vaginal tissue. On the other hand, the surface charge of the liposomes also plays an important role in the internalization of the encapsulated drug. Generally, positively surface-charged (cationic) liposomes allow fusion with negatively surface-charged epithelial cells, enabling efficient intracellular drug delivery.⁸ Since all the tested liposomes in this study were negatively surface charged, it is assumed that they exhibited lower affinity toward HeLa cells than the free drug, which is supported by Sangaré and collaborators,⁵³ who demonstrated that cationic liposomes encapsulating doxycycline were 2-fold more effective than free doxycycline against *C. trachomatis* in vitro. Therefore, the efficacy of the

positively surface-charged AZI-liposomes against *C. trachomatis* would be worthy to evaluate in future investigations.

Considering that the impact of antimicrobials on chlamydial growth was evaluated by direct qPCR with unpurified cell lysates, we measured the impact of the unpurified cell lysates on the performance of qPCR (Figure 7). Namely, it should be excluded whether the tested samples (AZI-liposomes and free AZI) had a direct inhibitory or stimulatory effect on the qPCR, influencing the dose-response curves of the AZI samples (AZI-liposomes and free AZI). To test this possible effect, Chlamydia-specific qPCRs were performed in the presence and absence of AZI (AZI-liposomes and free AZI). For that purpose, *C. trachomatis*-infected HeLa cell lysates were mixed with uninfected but AZI-treated (AZI-liposomes or free AZI) HeLa cell lysates. The final concentration of AZI was the highest concentration applied in the chlamydial growth inhibition experiments. Before cell lysis, similar to the dose-response curve experiments, the medium was removed, and the cells were washed with PBS. As a control, *C. trachomatis*-infected HeLa cell lysates were mixed with uninfected and AZI untreated HeLa cell lysates. If the AZI samples had no impact on the qPCR, then the AZI-containing samples had Ct levels similar to the Ct levels of the control.

The results presented in Figure 7 show that the detected chlamydial genome concentrations (Ct levels) of AZI samples (liposomes and the free drug) and the sample without AZI (control) were not significantly different (*t*-test; $p > 0.05$). Consequently, qPCR measurement of

chlamydial growth inhibition was not influenced by the applied compounds.

Biocompatibility studies

The biocompatibility of AZI-liposomes (CL-3, PGL-2 and DPGL-2) was investigated in vitro to determine the possible cytotoxic effects of the liposome constituents and AZI toward human cervical cells (HeLa cells). It was reported that AZI can synergistically enhance the antiproliferative activity of vincristine in cancer cells.⁵⁴ Furthermore, recent findings demonstrated that free AZI was cytotoxic to human fibroblasts at a concentration of 16 µg/ml (49% viability) and human keratinocytes at 64 µg AZI/ml (55% viability), while entrapment of AZI in liposomes significantly decreased skin cell cytotoxicity.¹⁷

The performed studies showed that all AZI-liposomes were compatible with the HeLa cells (Figure 8A). Even at the highest tested concentration (39.25 µg/ml), the viability of the cells remained above 70%, which is considered a noncytotoxic threshold.⁵⁵ Comparison of the different types of AZI-liposomes proved CL-3 to be the most compatible, while the HeLa cells were more sensitive to both types of elastic liposomes, particularly DPGL-2, at the highest tested AZI concentration, which covered and exceeded the MIC₅₀ and IC₅₀ values for *E. coli* and significantly surpassed the MIC values for *C. trachomatis*.

Evaluation of the empty liposomes (without encapsulated AZI) at concentrations that corresponded to AZI-liposomes proved the biocompatibility of all tested liposomes (Figure 8B). Therefore, the slightly increased sensitivity of the HeLa cells toward elastic AZI-liposomes (PGL-2, DPGL-2) (Figure 8A) is thought to be a result of the increased AZI release from the elastic liposomes (Figure 2), facilitated by the presence of 10% propylene glycol in dispersions of PGL-2 and DPGL-2, in addition to 15% monoacyl phosphatidylcholine inside the bilayers of DPGL-2 (Table 1). These assumptions are supported by recent findings demonstrating the sensitivity of human keratinocytes and fibroblasts to deformable liposomes, which was also affected by increased drug release and the presence of an edge activator (sodium deoxycholate).¹⁷

Although monoacyl phospholipids (lysophospholipids) have been proven to contribute to cell cytotoxicity at concentrations >50 µg/ml,⁵⁶ in this study the concentration of monoacyl phosphatidylcholine in DPGL-2 was significantly lower (39 µg/ml at the highest tested concentration), and the exposure time was 24 h in comparison to 72 h.⁵⁶

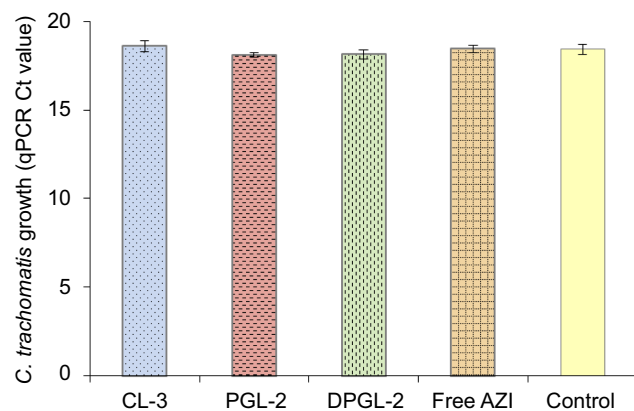


Figure 7 Estimation of the impact of AZI-liposomes and free AZI on qPCR.

Notes: Chlamydia-specific qPCRs were performed in the presence of AZI-liposomes and free AZI at the highest applied concentrations that were used in the dose-response curve measurement experiments. One-way ANOVA and Tukey's multiple comparison test, with $p < 0.05$ set as the minimal level of significance, were applied for comparison of qPCR Ct levels of CL-3, PGL-2, DPGL-2 and free AZI containing samples and control. No significant differences were found ($p > 0.05$). Data are presented as the mean Ct value \pm S.D. ($n=3$).

Abbreviations: AZI, azithromycin; Ct, cycle threshold; qPCR, quantitative polymerase chain reaction.

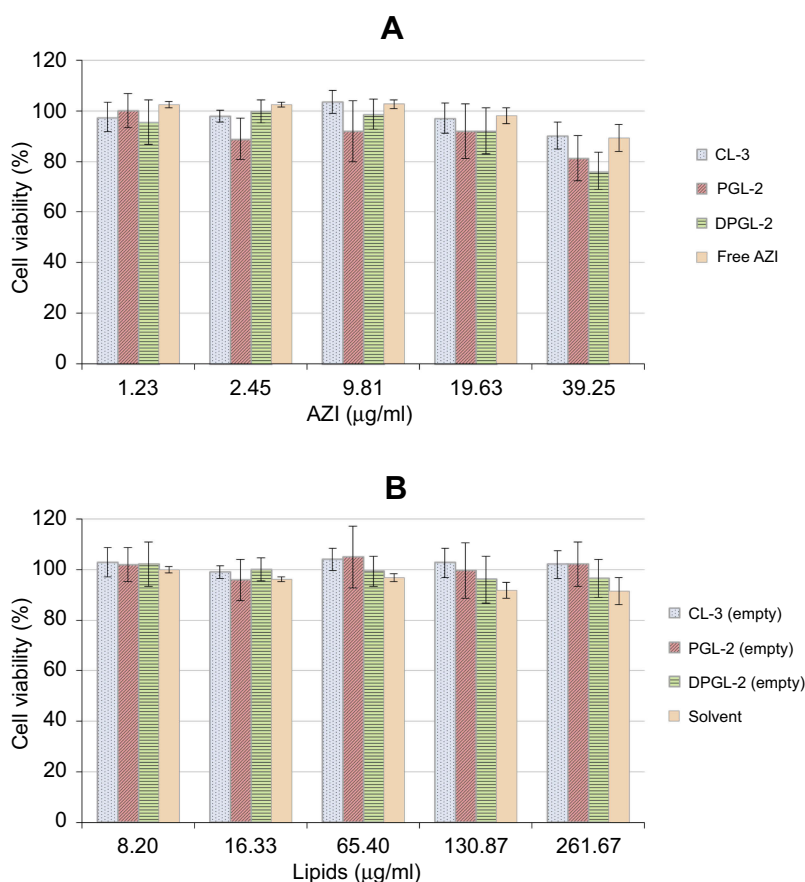


Figure 8 Viability of HeLa cells after incubation with different AZI-liposomes or free AZI (**A**) and with the corresponding empty liposomes (without AZI) or the solvent used to dissolve free AZI (**B**).

Note: The presented values are the mean \pm S.D. (n=3).

Abbreviation: AZI, azithromycin.

Hence, monoacyl phosphatidylcholine did not contribute to the cytotoxic effects of DPGL-2 under the applied conditions, even though there was a general trend of lower cell viability at higher AZI concentrations in DPGL-2 (Figure 8A). However, the possible cytotoxicity of DPGL-2 might be expected at higher concentrations of monoacyl phospholipid and with longer exposure of DPGL-2 to cervical cells (72 h).

The biocompatibility of the liposomes examined in this research could also be a consequence of their negative surface charge. Namely, cationic liposomes incorporating dimethyldioctadecyl ammonium bromide applied at higher concentrations significantly decreased the viability of A549 human lung cells and fibroblasts,^{17,57} in comparison to the corresponding negatively charged liposomes.

Conclusion

The results of all the performed studies indicate that targeting cervicovaginal infections requires careful tailoring of the

liposomal characteristics to yield stable liposomes that are biocompatible with cervical cells, allowing sustained AZI release and localization within vaginal tissue while enhancing antibacterial activity. The selection of AZI-liposomes should be based on the targeted bacterial strain. CL-3 would be optimal for preventing *E. coli* 700928 and K12 biofilms, while PGL-2 would be optimal for inhibiting the formation of *E. coli* 8739 biofilms. DPGL-2 exhibited the strongest potential for the eradication of *E. coli* biofilms and was the most promising among the examined liposomes toward *C. trachomatis*. Further formulation development should focus on ensuring the prolonged residence time of AZI-liposomes in the vaginal cavity by inclusion in a secondary vehicle, such as a mucoadhesive hydrogel.

Abbreviation list

AZI, azithromycin; CFU, colony forming units; CLs, conventional liposomes; Ct, cycle threshold; DPGLs, deformable propylene glycol liposomes; E, degree of liposome membrane

elasticity; EPC, egg phosphatidylcholine; EPG, egg phosphatidylglycerol sodium salt; IC₅₀, half-maximal inhibitory concentration; J, mass of extruded liposomes in elasticity measurements; LB, Luria-Bertani; MIC, minimum inhibitory concentration; MIC₅₀, concentration of the antibiotic that inhibited bacterial growth by 50%; MRSA, methicillin-resistant *Staphylococcus aureus*; PBS, phosphate-buffered saline; PCS, photon correlation spectroscopy; PDI, polydispersity index; PGLs, propylene glycol liposomes; qPCR, quantitative polymerase chain reaction; r_p, pore size of the membrane in elasticity measurements; r_v, average diameter of the liposomes after passage through a defined membrane in elasticity measurements; SLPC-80, monoacyl phosphatidylcholine from soybean; SPC-3, hydrogenated soybean phosphatidylcholine; VFS, vaginal fluid simulant.

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Disclosure

Dr Adyary Fallarero is currently employed by Thermo Fisher Scientific in Vantaa, Finland. The authors report no other conflicts of interest in this work.

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